

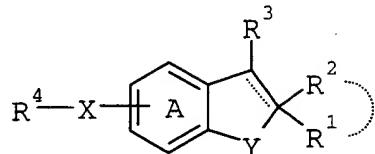
In the Claims

Please substitute the following claims 11, 12, 26 and 28 for the claims 11, 12, 26 and 28 now pending in the above-identified application.

Please cancel claims 22 and 25 without prejudice to the filing of future continuing applications.

Claims 1-10 (Cancelled)

11. (Currently Amended) A compound of the formula:



wherein R¹ and R² are each a C₁₋₆ alkyl or R¹ and R² form, taken together with the adjacent carbon atom, a piperidine optionally substituted by 1 to 3 substituents selected from the group consisting of C₁₋₆ alkyl, C₆₋₁₄ aryl and C₇₋₁₆ aralkyl; R³ is a phenyl optionally substituted by 1 to 3 substituents selected from the group consisting of halogen atoms, C₁₋₆ alkyl, C₁₋₆ alkoxy, amino, mono-C₁₋₆ alkylamino and di-C₁₋₆ alkylamino;

R⁴ is

(i) C₁₋₆ alkyl substituted by a phenyl or pyridyl, each of which is optionally substituted by 1 to 3 substituents selected from the group consisting of halogen atoms, C₁₋₆ alkyl, C₁₋₆ alkoxy, hydroxy, amino, mono-C₁₋₆ alkylamino, di-C₁₋₆ alkylamino and carboxy, or

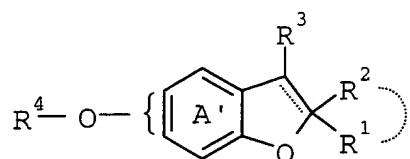
(ii) an acyl of the formula: -(C=O)-R^{5'} wherein R^{5'} is a phenyl or phenyl-C₁₋₆ alkyl, each of which is optionally substituted by 1 to 3 substituents selected from the group consisting of halogen atoms, C₁₋₆ alkyl, C₁₋₆ alkoxy, hydroxy, amino, mono-C₁₋₆ alkylamino, di-C₁₋₆ alkylamino and carboxy;

X is an oxygen atom;

Y is an oxygen atom; and

ring A is a benzene ring which is optionally further substituted by 1 to 3 substituents selected from the group consisting of halogen atoms, halogenated or unhalogenated C₁₋₆ alkyl, halogenated or unhalogenated C₁₋₆ alkoxy, amino, mono-C₁₋₆ alkylamino and di-C₁₋₆ alkylamino,
provided that when ----- is a single bond, R⁴ is not acyl,
and salts thereof.

12. (Currently Amended) A compound of the formula:



wherein R¹ and R² are each C₁₋₆ alkyl or R¹ and R² form, taken together with the adjacent carbon atom, a piperidine substituted by a C₁₋₆ alkyl or a C₇₋₁₆ aralkyl; R³ is a phenyl optionally substituted by 1 to 3 substituents selected from the group consisting of (1) C₁₋₆ alkyl, (2) di-C₁₋₆ alkylamino and (3) 6-membered saturated cyclic amino optionally substituted by a C₁₋₆ alkyl.

\mathbb{R}^4 is

- (i) a phenyl optionally substituted by 1 to 3 substituents selected from the group consisting of nitro and C_{1-6} alkyl-carboxamido,
- (ii) a C_{1-6} alkyl or C_{2-6} alkenyl group substituted by 1 to 3 of phenyl, quinolyl or pyridyl, each of which is optionally substituted by 1 to 3 substituents selected from the group consisting of C_{1-6} alkoxy, C_{1-6} alkylthio, C_{1-6} alkoxy-carbonyl, C_{1-6} alkylsulfonyl and C_{1-6} alkylsulfinyl, which C_{1-6} alkyl or C_{2-6} alkenyl group is optionally further substituted by a phenyl, carboxy or C_{1-6} alkoxy-carbonyl, or

(iii) an acyl of the formula: -(C=O)-R^{5"}

wherein R^{5"} is phenyl substituted by a C₁₋₆ alkoxy; and

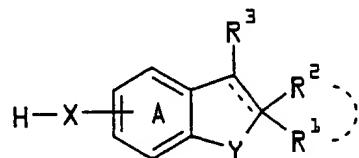
ring A' is a benzene ring which is optionally further substituted by 1 to 3 C₁₋₆ alkyl,

provided that when —— is a single bond, R⁴ is not acyl,

and salts thereof.

13. (Previously Presented) 3-(4-isopropylphenyl)-2,4,6,7-tetramethylbenzofuran-5-yl 4-methoxybenzoate, 3-(4-isopropylphenyl)-5-(4-methoxybenzyloxy)-2,4,6,7-tetramethylbenzofuran, 3-(4-isopropylphenyl)-5-(4-methoxybenzyloxy)-1',4,6,7-tetramethylspiro(benzofuran-2(3H), 4'-piperidine), or a salt thereof.

14. (Previously Presented) A process for producing a compound of Claim 11, which comprises reacting a compound of the formula:



wherein each symbol is as defined in Claim 11, or a salt thereof with a compound of the formula: R⁴-L wherein L represents a leaving group and R⁴ is as defined in Claim 11, or salt thereof.

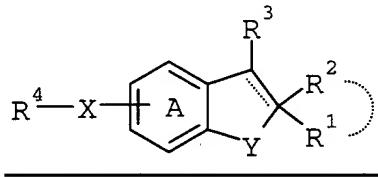
15. (Previously Presented) A pharmaceutical composition which comprises a compound of Claim 11, and a pharmaceutically acceptable carrier, excipient or diluent.

Claims 16-23 (Cancelled)

24. (Previously Presented) 3-(4-Isopropylphenyl)-5-(4-methoxybenzyloxy)-2,2,4,6,7-pentamethyl-2,3-dihydrobenzofuran.

25. (Cancelled)

26. (Currently Amended) A method of ~~claim 25, which is a method~~ for treating Alzheimer's disease in a mammal comprising administering to said mammal an effective amount of a compound of the formula



wherein R¹ and R² each represent an acyclic hydrocarbon group or a cycloalkyl group;

R³ represents an unsubstituted or substituted phenyl group;

R⁴ represents an aliphatic hydrocarbon group substituted by an unsubstituted or substituted aromatic group, which aliphatic hydrocarbon group is optionally further substituted;

X and Y each represent an oxygen atom;

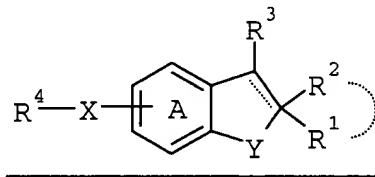
----- represents a single bond or a double bond;

and Ring A represents a benzene which is optionally further substituted apart from the group of the formula: -X-R⁴ wherein each symbol is as defined above,

or a salt thereof.

Claim 27 (Cancelled)

28. (Currently Amended) A method ~~of claim 22, which is a method~~ for treating Alzheimer's disease comprising administering to said mammal an effective amount of a compound of the formula:



wherein R¹ and R² are each a C₁₋₆ alkyl or R¹ and R² form, taken together with the adjacent carbon atom, a piperidine optionally substituted by 1 to 3 substituents selected from the group consisting of C₁₋₆ alkyl, C₆₋₁₄ aryl and C₇₋₁₆ aralkyl;

R³ is a phenyl optionally substituted by 1 to 3 substituents selected from the group consisting of halogen atoms, C₁₋₆ alkyl, C₁₋₆ alkoxy, amino, mono-C₁₋₆ alkylamino and di-C₁₋₆ alkylamino;

R⁴ is

(i) C₁₋₆ alkyl substituted by a phenyl or pyridyl, each of which is optionally substituted by 1 to 3 substituents selected from the group consisting of halogen atoms, C₁₋₆ alkyl, C₁₋₆ alkoxy, hydroxy, amino, mono-C₁₋₆ alkylamino, di-C₁₋₆ alkylamino and carboxy, or

(ii) an acyl of the formula: -(C=O)-R^{5'} wherein R^{5'} is a phenyl or phenyl-C₁₋₆ alkyl, each of which is optionally substituted by 1 to 3 substituents selected

from the group consisting of halogen atoms, C₁₋₆ alkyl, C₁₋₆ alkoxy, hydroxy,
amino, mono-C₁₋₆ alkylamino, di-C₁₋₆ alkylamino and carboxy;

X is an oxygen atom;

Y is an oxygen atom; and

ring A is a benzene ring which is optionally further substituted by 1 to 3

substituents selected from the group consisting of halogen atoms,

halogenated or unhalogenated C₁₋₆ alkyl, halogenated or unhalogenated C₁₋₆

alkoxy, amino, mono-C₁₋₆ alkylamino and di-C₁₋₆ alkylamino,

provided that when ----- is a single bond, R⁴ is not acyl,

and salts thereof.

Claim 29 (Cancelled)